## IN THE CLAIMS

Please amend the claims as follows:

- 1. (Withdrawn) A method of treating or preventing diseases of the eye, comprising, administering intraocularly a gene delivery vector which directs the expression of a neurotrophic factor, such that said disease of the eye is treated or prevented.
- 2. (Withdrawn) The method according to claim 1 wherein said neurotrophic factor is NGF, BDNF, CNTF, NT-3, or, NT-4.
- 3. (Withdrawn) The method according to claim 1 wherein said neurotrophic factor is a FGF.
- 4. (Withdrawn) The method according to claim 3 wherein said FGF is FGF-2, FGF-18, FGF-20, or, FGF-21.
- 5. (Withdrawn) The method according to claim 1 wherein said disease of the eye is macular degeneration.
- 6. (Withdrawn) The method according to claim 1 wherein said disease of the eye is diabetic retinopathy.
- 7. (Withdrawn) The method according to claim 1 wherein said disease of the eye is an inherited retinal degeneration.
- 8. (Withdrawn) The method according to claim 7 wherein said inherited retinal degeneration is retinitis pigmentosa.
- 9. (Withdrawn) The method according to claim 1 wherein said disease of the eye is glaucoma.

- 10. (Withdrawn) The method according to claim 1 wherein said disease of the eye is a surgery-induced retinopathy.
- 11. (Withdrawn) The method according to claim 1 wherein said disease of the eye is retinal detachment.
- 12. (Withdrawn) The method according to claim 1 wherein said disease of the eye is a photic retinopathy.
- 13. (Withdrawn) The method according to claim 1 wherein said disease of the eye is a toxic retinopathy.
- 14. (Withdrawn) The method according to claim 1 wherein said disease of the eye is a trauma-induced retinopathy.
- 15. (Withdrawn) The method according to claim 1 wherein said gene delivery vector is a retrovirus selected from the group consisting of HIV and FIV.
- 16. (Withdrawn) The method according to claim 1 wherein said gene delivery vector is a recombinant adeno-associated viral vector.
- 17. (Currently Amended) A method of inhibiting <u>angiogenesis in a</u>
  <u>diseased eye of a subject neovascular disease of the eye</u>, comprising, administering intraocularly a <u>retroviral</u> gene delivery vector which directs the expression of an anti angiogenic factor, such that <u>administration of said vector inhibits</u>
  <u>neovascularization of the diseased eye</u> said neovascular disease of the eye is inhibited.

- 18. (Currently Amended) The method according to claim 17 wherein said antiangiogenic factor is soluble Flt-1, PEDF, soluble Tie-2 receptor, or, a single chain anti-VEGF antibody.
- 19. (Currently Amended) The method according to claim 17 wherein said diseased neovascular disease of the eye is in a subject having diabetic retinopathy, wet AMD and or retinopathy of prematurity.
- 20. (Currently Amended) The method according to claim 17 wherein said retroviral gene delivery vector is a retrovirus selected from the group consisting of HIV and or FIV.
- 21. (Currently Amended) The method according to claim 17 wherein said retroviral gene delivery vector is a recombinant adeno-associated viral vector.
- 22. (Currently Amended) A <u>retroviral</u> gene delivery vector which directs the expression of a neurotrophic factor, or an anti-angiogenic factor, <u>whereby the administration of the retroviral gene delivery vector expressing the neurotrophic or anti-angiogenic factor into a diseased mammal inhibits neovascularization of the diseased tissue.</u>
- 23. (Withdrawn) The gene delivery vector according to claim 22 wherein said neurotrophic factor is NGF, BDNF, CNTF, NT-3, or, NT-4.
- 24. (Withdrawn) The gene delivery vector according to claim 22 wherein said neurotrophic factor is a FGF.
- 25. (Withdrawn) The gene delivery vector according to claim 22 wherein said FGF is FGF-2, FGF-5, FGF-18, FGF-20, or, FGF-21.

- 26. **(Currently Amended)** The <u>retroviral</u> gene delivery vector according to claim 22 wherein said anti angiogenic factor is soluble Flt-1, PEDF, soluble Tie-2 receptor, or, a single-chain anti-VEGF antibody.
- 27. (Currently Amended) The <u>retroviral</u> gene delivery vector according to claim 22 wherein said vector is generated from <u>a retrovirus</u> <u>an adeno-associated</u> <u>retrovirus or an alpha virus</u>.
- 28. (Currently Amended) The <u>retroviral</u> gene delivery vector according to claim 27 wherein the retrovirus is HIV or FIV.
  - 29. (Canceled)
- 30. (Withdrawn) A non-human animal model of neovascularization of the eye, comprising an animal having an angiogenic transgene in the eye.
- 31. (Withdrawn) The non-human animal model according to claim 30 wherein said neovascularization is retinal neovascularization.
- 32. (Withdrawn) The non-human animal model according to claim 30 wherein said neovascularization is choroidal neovascularization.
- 33. (Withdrawn) The non-human animal model according to claim 30 wherein said animal is a mouse or rat.
- 34. (Withdrawn) The non-human animal model according to claim 30 wherein said angiogenic transgene encodes VEGF.
- 35. (Withdrawn) The non-human animal model according to claim 30 wherein said angiogenic transgene encodes an angiopoietin.

- 36. (Withdrawn) A method for making a non-human animal model of neovascularization of the eye, comprising administering to a non-human animal a gene delivery vector which directs the expression of an angiogenic transgene.
- 37. (Withdrawn) The method according to claim 36 wherein said gene delivery vector is administered subretinally.
- 38. (Withdrawn) The method according to claim 36 wherein said gene delivery vector is administered intravitreally.
- 39. (Withdrawn) The method according to claim 36 wherein said gene delivery vector is rAV or rAAV.
- 40. (Withdrawn) The method according to claim 36 wherein said angiogenic transgene is a nucleic acid molecule which encodes VEGF.
- 41. (Withdrawn) The method according to claim 36 wherein said angiogenic transgene is a nucleic acid molecule which encodes an angiopoietin.
- 42. (Withdrawn) A method for determining the ability of an anti-angiogenic factor to inhibit neovascularization of the eye, comprising: (a) administering to an animal model according to any one of claims 30 to 35 an anti-angiogenic factor, and (b) determining the ability of said anti-angiogenic factor to inhibit neovascularization of the eye.
- 43. (Withdrawn) The method according to claim 42 wherein said anti-angiogenic factor is administered subretinally.
- 44. (Withdrawn) The method according to claim 42 wherein said anti-angiogenic factor is administered intravitreally.